

## Ch 10 Notes G.ink

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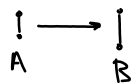
Types of Motion

Translational Motion

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Types of Motion

Translational Motion



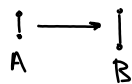
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Types of Motion

Translational Motion



Rotational Motion

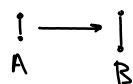
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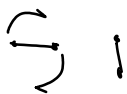
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Types of Motion

Translational Motion



Rotational Motion



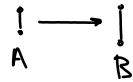
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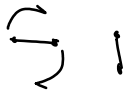
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Types of Motion

Translational Motion



Rotational Motion



Vibrational Motion

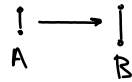
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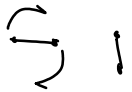
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Types of Motion

Translational Motion



Rotational Motion



Vibrational Motion



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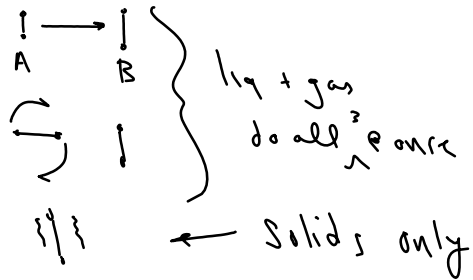
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### Types of Motion

Translational Motion

Rotational Motion

Vibrational Motion





Gases -

↳ Ideal Gases → conform to the Kinetic Molecular Theory of gases

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KMT

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Real Gases  
Deviate from ideal  
Behavior

### KMT

1. Samples of a gas consist of large #s of tiny particles
2. particles are in a const state of motion
3. particles undergo elastic collisions (No KE lost)
- ~~4~~ No forces of attraction B/w particles
- 5 KE of particles Directly prop to Kelvin T.

### Real Gases

Deviate from ideal Behavior

- Polar Gas molecules attract.
- Gases @ high P + low T

### Basic Gas Props

Expandable/compressible

Diffusion - spreading out of a gas  
in the absence of air currents

Effusion - passing of a gas molecule through  
an opening.

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↳ B/c large  $V$ ,  $D$  is low

unit:  $\frac{g}{L}$

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Expandable/compressible

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Effusion - passing of a gas molecule through  
an opening.

### Low Density

↳ B/c large  $V$ ,  $D$  is low

unit:  $\frac{g}{L}$

Fluidity - molecules  
can glide past each other  
(pourable)

Qualitative Description  
of Ideal Gas Behavior

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of Ideal Gas Behavior  
Variables

Qualitative Description  
of Ideal Gas Behavior

4 variables

Volume ( $V$ ) - space a gas  
occupies



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### Qualitative Description of Ideal Gas Behavior

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### Qualitative Description of Ideal Gas Behavior

#### 4 variables

Volume ( $V$ ) - space a gas occupies

Temperature ( $T$ ) - Avg KE of gas molecules  
(how fast they go)  
Always use K in calc

Moles ( $n$ ) - number of particles

Pressure ( $P$ ) - # of collisions  
gas molecules  
make w/ container  
walls

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$$\underline{p \propto T \text{ } p \text{ const } n + V}$$

$P \propto T$  if  $n$  &  $V$  const

As  $T \uparrow$ ,  $P \uparrow$  b/c as  $T \uparrow$   
particles move faster & faster  
so # of collisions  $\uparrow$  ;  $P \uparrow$



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$P$  vs  $T$  @ const  $n$  +  $V$

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$V$  vs.  $P$  @ const  $n$  +  $T$

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as  $V \uparrow$ ,  $P \downarrow$  b/c in a larger  
space there are less collisions  
B/w gas molec. so  $P \downarrow$

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$V$  vs  $T$  @ const  $n$  +  $P$

$T \uparrow$ ,  $V \uparrow$  b/c as molecules  
move faster they spread out  
occupying a larger  $V$ .

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$T \uparrow$ ,  $V \uparrow$  b/c as molecules  
move faster they spread out  
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Vice versa to all

$$\underline{P \propto n \text{ @ const } T + V}$$

If  $n \uparrow$ ,  $P \uparrow$  b/c more  
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$$\underline{n \propto V \text{ @ const } T + P}$$

$P \text{ vs } n \text{ @ const } T + V$

If  $n \uparrow$ ,  $P \uparrow$  b/c more particles in the same (fixed  $V$ ) mean more collisions

$n \text{ vs } V \text{ @ const } T + P$

If  $n \uparrow$ ,  $V \uparrow$  b/c more gas particles occupy more space.

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Pressure



Pressure   Measurements

$$P = \frac{\text{force}}{\text{area}}$$

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for gases  $P$  is collisions  
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$$\text{Standard } P = 760. \text{ mmHg} = 760. \text{ torr} = 1.00 \text{ atm} = 101.3 \text{ kPa}$$

(atmosphere)      (kilopascals)

1643 Evangelista Torricelli

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Measured atmospheric P  
using a Hg Barometer

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Measured atmospheric P

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STP - standard T + P  
0.0°C  
273 K  
1 atm, etc

Gas Law's  
Robert Boyle  
1662

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Gas Law's

Robert Boyle worked w/ V + P  
1662



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Gas Law's

Robert Boyle worked w/  $V + P$

1662

Saw if  $P$  doubled,  $V \downarrow$  by half

Inverse prop B/w  $V$  +  $P$

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$$y = \frac{k}{x}$$

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$$V = \frac{k}{P} \quad @ \text{ const } T + n$$

Inverse prop B/w  $V$  &  $P$

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$$VP = k$$

Inverse prop B/w  $V$  &  $P$

$$y = \frac{k}{x}$$

$$V = \frac{k}{P}$$

@ const  $T$  &  $n$

$$V_i P_i = k$$

Inverse prop B/w  $V$  &  $P$

$$y = \frac{k}{x}$$

$$V = \frac{k}{P}$$

@ const  $T + n$

$$V_1 P_1 = k$$

$V_2$

Inverse prop B/w  $V$  &  $P$

$$y = \frac{k}{x}$$

$$V = \frac{k}{P}$$

@ const  $T$  &  $n$

$$V_1 P_1 = k$$

$$V_2 P_2 = k$$



Inverse prop B/w  $V$  &  $P$

$$y = \frac{k}{x}$$

$$V = \frac{k}{P}$$

@ const  $T + n$

$$V_1 P_1 = k$$

$$V_2 P_2 = k$$

$$V_1 P_1 + V_2 P_2$$

@ const  $T + n$

Inverse prop B/w  $V$  &  $P$

$$y = \frac{k}{x}$$

$$V = \frac{k}{P} \quad @ \text{ const } T + n$$

$$V_1 P_1 = k$$

$$V_2 P_2 = k$$

$V_1 P_1 = V_2 P_2$   
@ const  $T + n$   
Boyle's Law

If 37.2 mL of  
a gas @ 24.0°C has  
its volume changed  
to 52.3 mL, what will  
the new P be if the  
original P was 682 torr  
and T remains const?

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$$V_1 P_1 = V_2 P_2$$

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$$V_1 P_1 = V_2 P_2$$

$V_1$

$P_1$

$V_2$

$P_2$

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$$V_1 P_1 = V_2 P_2$$

$$V_1 = 37.2 \text{ mL}$$

$$P_1 = 682 \text{ torr}$$

$$V_2 = 52.3 \text{ mL}$$

$$P_2 = ?$$

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$$V_2 = 52.3 \text{ mL}$$

$$P_2 = ?$$

$$P_2 = \frac{V_1 P_1}{V_2}$$



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$$V_2 = 52.3 \text{ mL}$$

$$P_2 = ?$$

$$P_2 = \frac{V_1 P_1}{V_2}$$

$$= \frac{(37.2 \text{ mL})(682 \text{ torr})}{52.3 \text{ mL}}$$

If 37.2 mL of  
a gas @ 24.0°C has  
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$$P_2 = \frac{V_1 P_1}{V_2}$$

$$= \frac{(37.2 \text{ mL})(682 \text{ torr})}{52.3 \text{ mL}}$$

$$= \boxed{485 \text{ torr}}$$

If 12.3 mL of He  
have a  $P$  of 94.5 kPa,  
what is  $V$  of the He  
when the  $P$  is 903 mmHg?

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$$V_1 = 12.3 \text{ mL} \quad V_2 = ?$$

$$P_1 = 94.5 \text{ kPa} \quad P_2 = 903 \text{ mmHg}$$

If 12.3 mL of He  
have a P of 94.5 kPa,  
what is V of the He  
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$$V_1 = 12.3 \text{ mL} \quad V_2 = ?$$

$$P_1 = 94.5 \text{ kPa} \quad P_2 = 903 \text{ mmHg} \times \frac{101.3 \text{ kPa}}{760 \text{ mmHg}} = 120.4 \text{ kPa}$$

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$$V_2 = \frac{V_1 P_1}{P_2} =$$

If 12.3 mL of He  
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when the P is 903 mmHg?  
 $V_1 = 12.3 \text{ mL}$   $V_2 = ?$

$$V_2 = \frac{V_1 P_1}{P_2} = \frac{(12.3 \text{ mL})(94.5 \text{ kPa})}{(120.4 \text{ kPa})}$$
$$= \boxed{9.65 \text{ mL}}$$

$$P_1 = 94.5 \text{ kPa} \quad P_2 = 903 \text{ mmHg} \times \frac{101.3 \text{ kPa}}{760 \text{ mmHg}} = 120.4 \text{ kPa}$$



1787 Jacques Charles  
worked w/ V+T

1787 Jacques Charles  
worked w/  $V \propto T$

Increased  $T$  by 1,  $V$  increased by  $\frac{1}{273}$

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Increased  $T$  by 1,  $V$  increased by  $\frac{1}{273}$   
Direct Prop!  
 $y = kx \rightarrow V_1 = kT_1, c \text{ const } n + P$

1787 Jacques Charles  
worked w/  $V$  +  $T$

Increased  $T$  by 1,  $V$  increased by  $\frac{1}{273}$   
Direct Prop!

$$y = kx \rightarrow V_1 = k T_1 \text{ @ const } n + P$$

$$V_2 = k T_2$$



1787 Jacques Charles  
worked w/ V + T

Increased T by 1, V increased by  $\frac{1}{273}$   
Direct Prop!

$$y = kx \rightarrow V_1 = k T_1 \text{ @ const } n + P$$
$$V_2 = k T_2 \quad \frac{V_1}{T_1} = k = \frac{V_2}{T_2}$$

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Increased T by 1, V increased by  $\frac{1}{273}$

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$$y = kx \rightarrow V_1 = k T_1 \text{ @ const } n + P \quad \frac{V_1}{T_1} = \frac{V_2}{T_2}$$
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Increased T by 1, V increased by  $\frac{1}{273}$

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Charles' Law

T must be K

A sample of  $\text{O}_2$  has a  
vol of 24.2 mL @ 57.3°C,  
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$$V_1 = 24.2 \text{ mL} \quad V_2 = 30.4 \text{ mL}$$

$$T_1 = 57.3^\circ\text{C} \quad T_2 = ?$$

A sample of  $\text{O}_2$  has a  
vol of 24.2 mL @ 57.3°C,  
what T is the gas when  
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space?

$$V_1 = 24.2 \text{ mL} \quad V_2 = 30.4 \text{ mL}$$

$$T_1: 57.3^\circ \text{C} \quad T_2 = ?$$
$$= 330.3 \text{ K}$$

A sample of  $\text{O}_2$  has a  
vol of 24.2 mL @ 57.3°C,  
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it occupies 30.4 mL of  
space?

$$V_1 = 24.2 \text{ mL} \quad V_2 = 30.4 \text{ mL}$$

$$T_1: 57.3^\circ \text{C} \quad T_2 = ?$$
$$= 330.3 \text{ K}$$

$$T_2 = \frac{V_2 T_1}{V_1}$$



A sample of  $\text{C}_3\text{H}_8$  has a  
vol of 24.2 mL @ 57.3°C,  
what T is the gas when  
it occupies 30.4 mL of  
space?

$$V_1 = 24.2 \text{ mL} \quad V_2 = 30.4 \text{ mL}$$

$$T_1: 57.3^\circ \text{C} \rightarrow T_2 = ? \\ = 330.3 \text{ K}$$

$$T_2 = \frac{V_2 T_1}{V_1} = \frac{(30.4 \text{ mL})(330.3 \text{ K})}{(24.2 \text{ mL})}$$

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$$T_2 = \frac{V_2 T_1}{V_1} = \frac{(30.4 \text{ mL})(330.3 \text{ K})}{(24.2 \text{ mL})} \\ = 415 \text{ K}$$

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$$V_1 = 24.2 \text{ mL} \quad V_2 = 30.4 \text{ mL}$$

$$T_1: 57.3^\circ (+273) \quad T_2 = ? \\ = 330.3 \text{ K}$$

$$T_2 = \frac{V_2 T_1}{V_1} = \frac{(30.4 \text{ mL})(330.3 \text{ K})}{(24.2 \text{ mL})} \\ = 415 \text{ K} - 273 = \boxed{142^\circ \text{C}}$$

1808 Joseph Gay-Lussac

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P vs T - If  $T \uparrow$  by 1,  $P \uparrow$  by  $\frac{1}{273}$

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1808 Joseph Gay-Lussac

P vs T - If  $T \uparrow$  by 1,  $P \uparrow$  by  $\frac{1}{273}$

Direct Prop -  $y = kx$

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$$P = kT$$

$$\frac{P}{T} = k$$



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or

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or

$P_1 T_2 = P_2 T_1$  Gay-Lussac's,  
T in Kelvin Law

Boyle's  $V_1 P_1 = V_2 P_2$

Charles's  $V_1 T_2 = V_2 T_1$

Gay-Lussac's  $P_1 T_2 = P_2 T_1$

$$\left. \begin{array}{l} \text{Boyle's} \quad V_1 P_1 = V_2 P_2 \\ \text{Charles's} \quad V_1 T_2 = V_2 T_1 \\ \text{Gay-Lussac's} \quad P_1 T_2 = P_2 T_1 \end{array} \right\} \begin{array}{l} V_1 P_1 T_2 = V_2 P_2 T_1 \\ \text{Combined Gas Law} \end{array}$$

A sample of  $\text{NO}_2$   
has a vol of 24.5 mL @  
7.62 atm and 30.4°C. What  
will the Vol of the gas be  
@ STP?

$$V_1 P_1 T_2 = V_2 P_2 T_1$$

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will the vol of the gas be

@ STP?

$$V_1 = 24.5 \text{ mL} \quad V_2 = ?$$

$$P_1 = 7.62 \text{ atm} \quad P_2 = 1.00 \text{ atm}$$

$$T_1 = 30.4^\circ\text{C} \rightarrow 303.4 \text{ K} \quad T_2 = 0^\circ\text{C} = 273 \text{ K}$$

$$V_1 P_1 T_2 = V_2 P_2 T_1$$



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$$T_1 = 30.4^\circ\text{C} \rightarrow 303.4 \text{ K} \quad T_2 = 0^\circ\text{C} \rightarrow 273 \text{ K}$$

$$V_1 P_1 T_2 = V_2 P_2 T_1$$

$$V_2 = \frac{V_1 P_1 T_2}{P_2 T_1}$$

$$= \frac{(24.5 \text{ mL})(7.62 \text{ atm})(273 \text{ K})}{(1.00 \text{ atm})(303.4 \text{ K})}$$

=

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$$V_1 P_1 T_2 = V_2 P_2 T_1$$

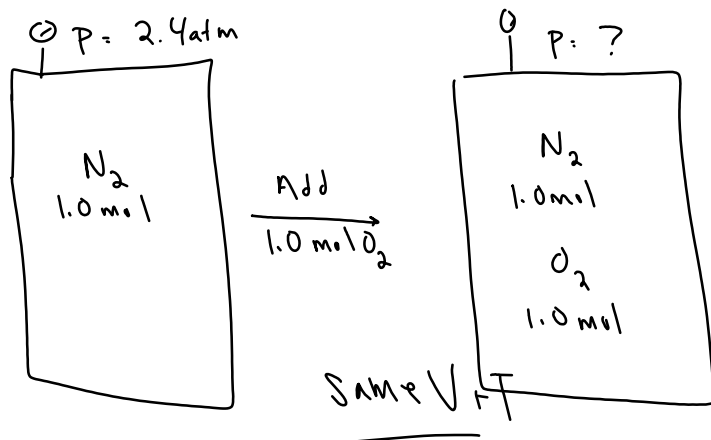
$$V_2 = \frac{V_1 P_1 T_2}{P_2 T_1}$$

$$= \frac{(24.5 \text{ mL})(7.62 \text{ atm})(273 \text{ K})}{(1.00 \text{ atm})(303.4 \text{ K})}$$

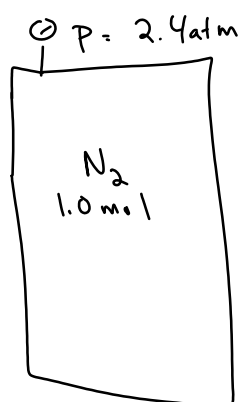
$$= 168 \text{ mL}$$

Dalton's Law of Partial P

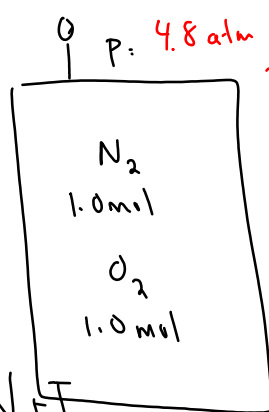
Dalton's Law of Partial P



Dalton's Law of Partial P



Add  
 $1.0 \text{ mol } O_2$



b/c double # of  
particles + P doubles

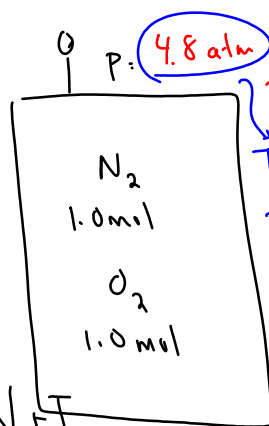
Same V & T

Dalton's Law of Partial P



Add  
 $\xrightarrow{1.0 \text{ mol } O_2}$

Same V & T



b/c double # of  
particles + P doubles

Total P ( $P_T$ )

$$P_T = P_{N_2} + P_{O_2}$$

Dalton's Law of Partial P

In a mixture of gases  
The total P of the sample  
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the partial P of each gas  
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Dalton's Law of Partial P

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The total P of the sample  
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$$P_T = P_1 + P_2 + P_3 \text{ etc}$$



### Dalton's Law of Partial P

In a mixture of gases  
The total P of the sample  
is equal to the sum of  
the partial P of each gas  
present

$$P_T = P_1 + P_2 + P_3 + \dots$$

$$P_T = 640 \text{ torr}$$

$$N_2 = 500 \text{ torr}$$

$$O_2 = 45 \text{ torr}$$

He: ?

### Dalton's Law of Partial P

In a mixture of gases  
The total P of the sample  
is equal to the sum of  
the partial P of each gas  
present

$$P_T = P_1 + P_2 + P_3 + \dots$$

$$P_T = 640 \text{ torr}$$

$$N_2 = 500 \text{ torr}$$

$$O_2 = 75 \text{ torr}$$

$$He = ?$$

$$P_T = P_{N_2} + P_{O_2} + P_{He}$$

### Dalton's Law of Partial P

In a mixture of gases  
The total P of the sample  
is equal to the sum of  
the partial P of each gas  
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$$P_T = P_1 + P_2 + P_3 + \dots$$

$$P_T = 640 \text{ torr}$$

$$N_2 = 500 \text{ torr}$$

$$O_2 = 45 \text{ torr}$$

$$He = ?$$

$$P_T = P_{N_2} + P_{O_2} + P_{He}$$

$$640 \text{ torr} = 500 \text{ torr} + 45 \text{ torr} = P_{He}$$

### Dalton's Law of Partial P

In a mixture of gases  
The total P of the sample  
is equal to the sum of  
the partial P of each gas  
present

$$P_T = P_1 + P_2 + P_3 + \dots$$

$$P_T = 640 \text{ torr}$$

$$N_2 = 500 \text{ torr}$$

$$O_2 = 95 \text{ torr}$$

$$He = ?$$

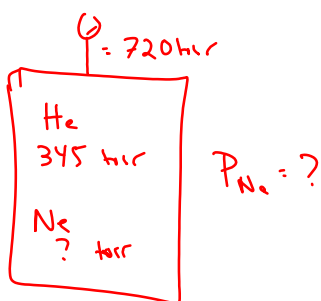
$$P_T = P_{N_2} + P_{O_2} + P_{He}$$

$$640 \text{ torr} = 500 \text{ torr} + 95 \text{ torr} = P_{He}$$

$$P_{He} = 95 \text{ torr}$$

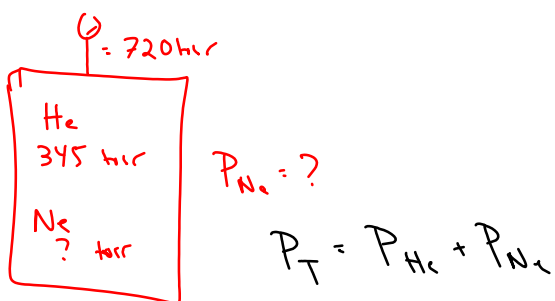
## Ch 10 Notes G.ink

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## Ch 10 Notes G.ink

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## Ch 10 Notes G.ink

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$P = 720 \text{ torr}$

He
345 torr
Ne
? torr

$P_{\text{Ne}} = ?$

$$P_T = P_{\text{He}} + P_{\text{Ne}}$$

$$720 \text{ torr} = 345 \text{ torr} + P_{\text{Ne}}$$

## Ch 10 Notes G.ink

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$P = 720 \text{ torr}$

He
345 torr
Ne
? torr

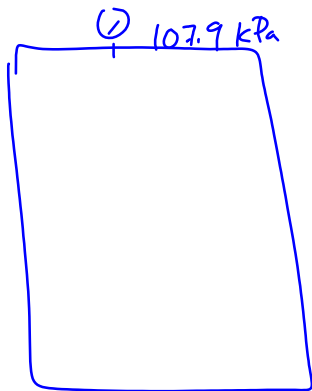
$P_{\text{Ne}} = ?$

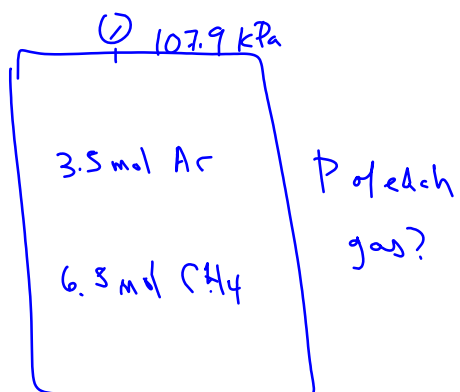
$$P_T = P_{\text{He}} + P_{\text{Ne}}$$

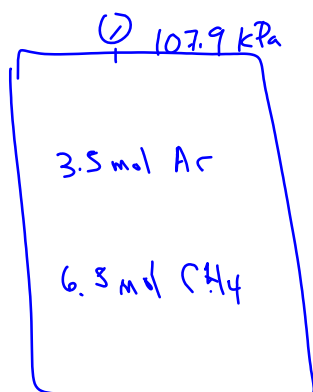
$$720 \text{ torr} = 345 \text{ torr} + P_{\text{Ne}}$$

$$P_{\text{Ne}} = 375 \text{ torr}$$



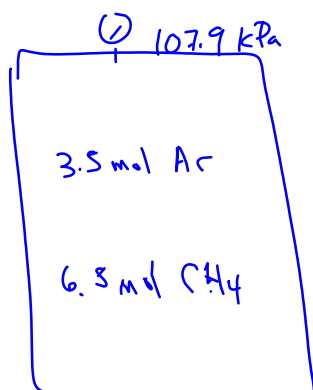






Partial  
gas?

$$P_{\text{Ar}} = \frac{3.5 \text{ mol}}{10.0 \text{ mol}} \times 107.9 \text{ kPa} = 37.8 \text{ kPa}$$



P of each  
gas?

$$P_{\text{Ar}} = \frac{3.5 \text{ mol}}{10.0 \text{ mol}} \times 107.9 \text{ kPa} = 37.8 \text{ kPa}$$

$$P_{\text{CH}_4} = 107.9 \text{ kPa} - 37.8 \text{ kPa} = 70.1 \text{ kPa}$$

Water Vapor Pressure (WVP,  $P_{wv}$ )

Water Vapor Pressure (WVP,  $P_{H_2O}$ )

When a gas is collected over water, some water will evaporate into the gas exerting additional P (WVP)

Now the collected gas is "wet" and we must adjust our calc's for the WVP.

If 23.2 mL of  $H_2$  gas is  
collected over  $H_2O$  @  $25^\circ C$  +  
738 torr of P, what is the  
V of the dry gas @ STP?

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 $V_1$  $P_1$  $T_1$  $V_2$  $P_2$  $T_2$

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V of the dry gas @ STP?

(wvp @  $25.00^\circ C = 23.8 \text{ mm}$ )

$$V_1 = 23.2 \text{ mL}$$

$$V_2 = ?$$

$$P_1$$

$$P_2 = 760 \text{ torr}$$

$$T_1 = 25.0^\circ C + 273 = 298 \text{ K} \quad T_2 = 273 \text{ K}$$

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(wvp @  $25.00^\circ C = 23.8 \text{ mmHg}$ )

$$V_1 = 23.2 \text{ mL} \quad V_2 = ?$$

$$P_1 = 738 \text{ torr} - 23.8 \text{ torr} = 714.2 \text{ torr} \quad P_2 = 760 \text{ torr}$$

$$T_1 = 25.0^\circ C + 273 = 298 \text{ K} \quad T_2 = 273 \text{ K}$$

## Ch 10 Notes G.ink

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$$V_2 = \frac{V_1 P_1 T_2}{P_2 T_1} = \frac{(23.2 \text{ mL})(714.2 \text{ torr})(273 \text{ K})}{(760 \text{ torr})(298 \text{ K})}$$

## Ch 10 Notes G.ink

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$$= \boxed{20.0 \text{ mL}}$$